Installation Report

The installation report is an important form for Arcoma to receive to keep track of delivered systems and their status. The report is required from all performed installations to comply with CFR 21 §1020.30. The CE-mark to MDD Class II products is fulfilled through MDD ANNEX II 93/42/EEC where our Quality system is an essential part.

The installation report forms are delivered with each system (included in the Service and Installation Manual). There is also a digital form (this document) available at our website (www.arcoma.se/contact).

Please send the report to service@arcoma.se.

Sending the report confirms that you have installed the unit and that it is working properly on site. If you encounter product related issues during the installation, it is important that we receive this information, please contact service@arcoma.se (+46 470 70 69 70).

INFORMATION FROM DEALER

Equipment type:
System serial number:
Date:
Distributor/Dealer/Partner:
Installer:

Hospital/address:
Department:
Lab/Room:

I confirm that this installation is performed in accordance with the installation chapter of Service & Installation Manual with rev. no:

Date:

Signature: .....

Other comments/input: